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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Group:	1626	}	
Confirmation No.:	9842	}	I hereby certify that this correspondence is being
		}	filed electronically in the United States Patent and
		}	Trademark Office
Application No.:	10/725,191	}	
		}	on <u>March 23, 2007</u>
Invention:	PHOSPHORAMIDE COMPOUNDS	}	
		}	(Signature)
Applicant:	Richard F. Borch et al.	}	<u>Charles L. Teyman</u>
		}	(Printed Name)
Filed:	December 1, 2003	}	
Attorney Docket:	3220-200986	}	
Examiner:	Golam M. Shameem	}	

SUBMISSION MADE WITH AN RCE:

Supplemental Information Disclosure Statement

Mail Stop RCE
Commissioner for Patents
P. O. Box 1450
Alexandria, Virginia 22313-1450

Sir:

Applicants have concurrently filed a request for continued examination (RCE) under 37 C.F.R. § 1.114 in a separate paper. In association with that RCE, Applicants hereby request entry and consideration of the instant Submission, which Submission includes a Supplemental Information Disclosure Statement (IDS), Form 1449, and all required copies of the references cited therein.

This IDS is filed in the application identified above pursuant to 37 C.F.R. § 1.56. No representation is intended that a complete search has been made of the prior art or that no better art references than the references cited in the IDS are available. The filing of this IDS shall not be construed to be an admission that the information cited in the IDS is, or

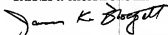
1.98(a)(2)(ii), copies of the cited U.S. patents are not provided herewith. Copies of the cited foreign patent documents and other references (non-patent) have been provided.

The cited references are not believed to disclose or suggest the invention recited in the claims of the above-identified application. It is therefore believed that the claimed invention is patentably distinguishable over the cited references.

Applicants have authorized, concurrent to filing the instant IDS, Barnes & Thornburg LLP Deposit Account 10-0435 to be debited in the amount of \$575.00, which amount is the sum of the \$180.00 fee set forth under 37 C.F.R. 1.17(p) for filing an Information Disclosure Statement, and the \$395.00 fee set forth under 37 C.F.R. § 1.17(e) for requesting continued examination under small-entity status.

Applicants do not believe that any additional fees are due for filing the instant Information Disclosure Statement. However, the Commissioner is hereby authorized to charge any additional fees that may be required, or credit any overpayment, to Barnes & Thornburg LLP Deposit Account No. 10-0435, with reference to our matter number 3220-200986.

Respectfully submitted,
BARNES & THORNBURG LLP



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Registration No. 48,480
Agent for Applicants

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317-231-7401

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE				ATTY. DOCKET NO.: 3220-200986		SERIAL NO.: 10/725,191	
INFORMATION DISCLOSURE STATEMENT				APPLICANT(S): Borch et al.		FILING DATE: Dec. 1, 2003	
INFORMATION DISCLOSURE STATEMENT				GROUP: 1626			

U.S. PATENT DOCUMENTS							
*Examiner Initial	Document Number	Date	Name	Class	Subclass	Filing Date if appropriate	
	AA	5,233,031					
	AB	6,903,081					
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	Document Number	Date	Country	Class	Subclass	Translation Yes No	
	AL	WO 93/06120	April 1, 1993	PCT			
	AM	WO 01/74527	Oct. 11, 2001	PCT			
	AN						
	AO						

OTHER REFERENCES (Including Author, Title, Date, Pertinent Pages, Etc.)

	AR	"The Synthesis of O-Monosaccharidyl-Methoxycarbonyl-Phosphoramidates by Arbuzov Reaction," Chen et al., Chinese Chemical Letters, vol. 6, No. 1, pp. 23-26, 1995
	AS	"Design and Synthesis of Lipophilic Phosphoramidate dAT-MP Prodrugs Expressing High Potency Against HIV in Cell Culture: Structural Determinants for <i>in vitro</i> Activity and QSAR," Silek et al., J. Med. Chem. 1999, 42, 4122-4128
	AT	"Phosphoramidates as Potent Prodrugs of Anti-HIV Nucleotides: Studies in the Amino Region," McGuigan et al., Antiviral Chemistry & Chemotherapy (1996) 7(1), 31-36
	AU	"Design and Synthesis of Novel Nucleotide Prodrugs," Meyers et al., American Association for Cancer Research 2000 (Abstract Book Page), Abstract #100710, dated December 2, 1999
	AV	"Protein Tyrosine Kinases and Cancer," Kolibaba et al., Biochimica et Biophysica Acta 1333 (1997) F217-F248
	AW	"Receptor Protein-Tyrosine Kinases and Their Signal Transduction Pathways," van der Geer et al., Annu. Rev. Cell Biol., 1994, 10:251-337
	AX	"Src Homology Region 2 Domains Direct Protein-Protein Interactions in Signal Transduction," Moran et al., Proc. Natl. Acad. Sci. USA, Vol. 87, Nov. 1990, pp. 8622-8626
	AY	"Binding of SH2 Domains of Phospholipase C γ 1, GAP, and Src to Activated Growth Factor Receptors," Anderson et al., Science, Vol. 250, Nov. 16, 1990, 979-982
	AZ	"Oncogenes and Signal Transduction," Cantley et al., Cell, Vol. 64, January 25, 1991, 281-302

Examiner	Date Considered
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*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 606.
 Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE				ATTY. DOCKET NO.: 3220-200986		SERIAL NO.: 10725,191	
INFORMATION DISCLOSURE STATEMENT				APPLICANT(S): Borch et al.		FILING DATE: Dec. 1, 2003	
				GROUP: 1828			

U.S. PATENT DOCUMENTS							
*Examiner Initial		Document Number	Date	Name	Class	Subclass	Filing Date if appropriate
	BA						
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FOREIGN PATENT DOCUMENTS							
		Document Number	Date	Country	Class	Subclass	Translation Yes No
	BL						
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BR	"Recognition and Specificity in Protein Tyrosine Kinase-Mediated Signalling," Songyang et al., TIBS 20, November 1995, 470-475
BS	"Targeting Signal Transduction in the Discovery of Antiproliferative Drugs," Saitel et al., Chemistry & Biology, 1998, Vol. 3, No. 11, 887-983
BT	"Peptide Inhibitors of src SH3-SH2-Phosphoprotein Interactions," Gilmer et al., The Journal of Biological Chemistry, Vol. 269, No. 50, December 16, 1994, 31711-31719
BU	"Structure-Based Design of a Novel Series of Nonpeptide Ligands That Bind to the pp60 ^{src} SH2 Domain," Lunney et al., J. Am. Chem. Soc. 1997, 119, 12471-12478
BV	"Design, Synthesis, and Co-crystal Structure of a Nonpeptide SrcSH2 Domain Ligand," Plummer et al., J. Med. Chem. 1997, 40, 3719-3725
BW	"Inhibition of SH2 Domain/Phosphoprotein Association by a Nonhydrolyzable Phosphonopeptide," Domchek et al., Biochemistry 1992, 31, 9865-9870
BX	"Nonhydrolyzable Phosphotyrosyl Mimetics for the Preparation of Phosphatase-resistant SH2 Domain Inhibitors," Burke, Jr., et al., Biochemistry, 1994, 33, 6490-6494
BY	L-O-(2-Malonyl)tyrosine: A New Phosphotyrosyl Mimetic for the Preparation of Src Homology 2 Domain Inhibitory Peptides," Ye et al., J. Med. Chem. 1995, 38, 4270-7275
BZ	"Conformationally Constrained Phosphotyrosyl Mimetics Designed as Monomeric Src Homology 2 Domain Inhibitors," Burke Jr., et al., J. Med. Chem. 1995, 38, 1389-1399

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INFORMATION DISCLOSURE STATEMENT				APPLICANT(S): Borch et al.			
				FILING DATE: Dec. 1, 2003		GROUP: 1626	

U.S. PATENT DOCUMENTS							
*Examiner Initial		Document Number	Date	Name	Class	Subclass	Filing Date if appropriate
	CA						
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	CI						
	CJ						
	CK						

FOREIGN PATENT DOCUMENTS							
		Document Number	Date	Country	Class	Subclass	Translation Yes No

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	CR "4'-O-[2-(2-Fluoromethyl)]-L-Tyrosine: A Phosphotyrosyl Mimic for the Preparation of Signal Transduction Inhibitory Peptides," Burke Jr., et al., J. Med. Chem. 1996, 39, 1021-1027
	CS "Phosphotyrosine-Containing Dipeptides as High-Affinity Ligands for the p56 ^{lck} SH2 Domain," Linas-Brunet et al., J. Med. Chem. 1999, 42, 722-729
	CT "Acquisition of High-Affinity, SH2-Targeted Ligands via a Spatially Focused Library," Lee et al., J. Med. Chem. 1999, 42, 784-787
	CU "Ligands for the Tyrosine Kinase p56 ^{lck} SH2 Domain: Discovery of Potent Dipeptide Derivatives with Monocharged, Nonhydrolyzable Phosphate Replacements," Beaulieu et al., J. Med. Chem. 1999, 42, 1757-1768
	CV "Synthesis and Biological Evaluation of 5-Fluoro-2'-Deoxyuridine Phosphoramidate Analogs," Fries et al., J. Med. Chem., 1995, 38, 2672-2680
	CW "Synthesis and Biological Activity of Novel 5-Fluoro-2'-Deoxyuridine Phosphoramidate Prodrugs," Meyers et al., J. Med. Chem., 2000, 43, 4313-4318
	CX "Synthesis and Evaluation of Nitroheterocyclic Phosphoramidates as Hypoxic-Selective Alkylating Agents," Borch et al., J. Med. Chem., 2000, 43, 2258-2265
	CY Gibbs et al., Current Medicinal Chemistry, Vol. 6, No. 12, October 2001, pp. 1437-1465
	CZ Borch et al., European Journal of Cancer. Supplement, Pergamon, Oxford, GB, Vol. 2, No. 8, September 2004, p. 178

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*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609.
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U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE INFORMATION DISCLOSURE STATEMENT				ATTY. DOCKET NO.: 3220-200996 APPLICANT(S): Borch et al. FILING DATE: Dec. 1, 2003		SERIAL NO.: 10725,191 GROUP: 1626	
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U.S. PATENT DOCUMENTS							
*Examiner Initial	Document Number	Date	Name	Class	Subclass	Filing Date if appropriate	
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FOREIGN PATENT DOCUMENTS							
Document Number	Date	Country	Class	Subclass	Translation Yes No		
DL							
DM							
DN							
DO							

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DR	Weidong et al., American Chemical Society, Abstracts of Papers at the National Meeting, ACS, Washington, D.C., Vol. 228, Part 1, 26 August 2004, p. U930	
DS		
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Examiner	Date Considered
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